

## Development of novel synNotch CART cell therapy in patients with recurrent EGFRvIII+ glioblastoma

### Grant Award Details

Development of novel synNotch CART cell therapy in patients with recurrent EGFRvIII+ glioblastoma

**Grant Type:** Therapeutic Translational Research Projects

**Grant Number:** TRAN1-12905

**Project Objective:** The objective of this project is to carry out the work necessary to have a successful pre-IND meeting in which the FDA agrees to the planned pivotal GLP safety studies to be done prior to IND filing for the product: EGFRvIII-synNotch primed EphA2/IL-13R $\alpha$ 2 CART cells (these are human T cells transduced with a lentiviral vector to express EGFRvIII-synNotch that, when bound to EGFRvIII, stimulates expression of anti-EphA2/IL-13R $\alpha$ 2 CAR). The work planned includes 1) process development for GMP compatible manufacturing of the product, 2) GMP manufacture of the viral vector needed to make the transduced T cells, 3) In vitro and in vivo assessments of function, phenotype, efficacy and biodistribution; 4) Pilot safety studies; 5) clinical protocol development; and 6) preparation of a pre-IND meeting package, submission to FDA, and holding the pre-IND meeting.

**Investigator:**

<b>Name:</b>	Hideho Okada
<b>Institution:</b>	University of California, San Francisco
<b>Type:</b>	PI

**Disease Focus:** Brain Cancer, Cancer, Solid Tumors

**Human Stem Cell Use:** Somatic Cell

**Award Value:** \$4,556,536

**Status:** Active

### Grant Application Details

**Application Title:** Development of novel synNotch CART cell therapy in patients with recurrent EGFRvIII+ glioblastoma

**Public Abstract:****Translational Candidate**

Human T cells transduced with a lentiviral vector encoding anti-EGFRvIII synNotch-primed anti-EphA2/IL-13R $\alpha$ 2 chimeric antigen receptor.

**Area of Impact**

Glioblastoma is the most common malignant brain tumor, affecting approximately 3 out of 100,000 people/year in the USA with extremely poor prognosis.

**Mechanism of Action**

In our proposed system, the first antigen EGFRvIII, which is expressed exclusively but heterogeneously on glioblastoma cells, primes the T cells to induce expression of a CAR that recognizes EphA2 and IL-13R $\alpha$ 2, thereby eradicating glioblastoma cells expressing either EphA2 or IL-13R $\alpha$ 2. Efficacy was long-lasting and superior to conventional CART cells. The superb efficacy of these synNotch-CART cells was associated with excellent persistence (>100 days in vivo) and T stem memory cell phenotype.

**Unmet Medical Need**

Glioblastoma is the most common malignant primary brain tumor, affecting approximately 3 out of 100,000 individuals/year in the USA. Despite surgical resection, radiation and chemotherapy, prognosis remains poor with a 100% recurrence rate and median overall survival of approximately 20 months.

**Project Objective**

Successful submission of a Pre-IND application

**Major Proposed Activities**

- Process development for manufacturing of EGFRvIII-primed EphA2/IL-13R $\alpha$ 2 CART cells
- In vivo (rodent) studies to determine preclinical efficacy and safety of the proposed cell products
- Development of the clinical trial protocol, consent form and clinical standard operating procedures (SOPs)

**Statement of Benefit to California:**

Because the current California's population is nearly 40 million, approximately 1,200 people are likely to be diagnosed with this devastating disease every year. The Brain Tumor Center within the Department of Neurosurgery at UCSF is one of the most established brain tumor research and treatment centers in the world. Our scientists and health care clinicians work in partnership to translate laboratory findings into new or improved forms of clinical therapy for patients in California.

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